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(54) Title: A METHOD OF PREPARING CROSS-LINKED ENZYME AGGREGATES

(57) Abstract: The invention relates to a method of preparing cross-linked enzyme aggregates (CLEAs) comprising the steps of a) precipitating a dissolved enzyme present in a liquid medium; b) cross-linking with a cross-linker, and c) optionally, washing the cross-linked enzyme aggregate. According to the invention at least one treatment is performed chosen from the group consisting of I) performing step a) in the presence of a compound selected from the group consisting of i) a crown ether, and ii) a surfactant, and II) performing step c) with a second liquid differing in organic composition from the liquid.

A method of preparing cross-linked enzyme aggregates

The present invention relates to a method of preparing cross-linked enzyme aggregates comprising the steps of
a) precipitating a dissolved enzyme present in a liquid medium;

5 b) cross-linking with a cross-linker, and

c) optionally, washing the cross-linked enzyme aggregate.

Such a method is generally known in the art. It is used to prepare enzyme preparations for performing various enzyme-catalysed reactions. For example, Cao L. et al. (Organic Letters vol. 2 (10) p. 1361-1364, 2000) describe the preparation of penicillin acylase-based aggregates. The cross-linked aggregates disclosed are less active than the native enzyme.

The object of the present invention is to provide a
15 method making it possible to achieve aggregates with increased enzymatic activity.

To this end, the invention is characterized in that at least one treatment is performed chosen from the group consisting of I) performing step a) in the presence of a compound selected from the group consisting of i) a crown ether, and ii) a surfactant, and II) performing step c) with a second liquid differing in organic composition from the liquid.

Applicants have surprisingly found that such a treatment may result in an cross-linked enzyme aggregate (CLEA) with an enhanced activity. While the present invention does neither guarantee this enhanced activity for a particular enzyme used to catalyse a particular reaction, nor allows for a prediction of an enhanced activity, it does provide for an extra parameter to be controlled to achieve a better aggregate. That is, for a particular enzyme, it may be investigated with little effort and using simple and routine experimentation whether its catalytic activity can be enhanced using the treatment. From a range of enzymes capable of catalyzing the desired reaction, the best CLEA can easily and 35 with little effort be selected. Because the CLEA is insoluble, it can be washed with a second liquid which is com-

pletely aqueous, organic, or a mixture of the two. After the treatment, the CLEA may be lyophilized or stored as a suspension for future use.

Overbeeke P.L.A. et al. Describe in J. of Mol. Cat. 5 B: Enzymatic 10, p. 385-393 (2000) a method in which a lipase is crystallized under the influence of PEG in the presence of 2-methyl-2,4-pentanediol, and crystals formed are cross-linked. In contrast, the present invention makes use of a precipitating agent to form aggregates instead of crystals.

10 Preferably, step a) and step b) are performed simultaneously.

This allows for a simplified procedure, requiring less time and effort.

According to a preferred embodiment, the compound 15 is removed after cross-linking.

It is not necessary that the surfactant and/or crown ether are present during the catalytic reaction to be performed by the CLEA to maintain the high activity. Thus the compound may be removed, in case it may interfere with the 20 reaction to be catalysed, or for other reasons. Removal may be achieved using any method known in the art, and in particular through dialysis or washing using centrifugation.

According to an important application of the method according to the invention, the enzyme is a lipase.

25 Hence, the invention also relates to a method for performing a lipase-catalysed reaction, said reaction being chosen from the group consisting of i) transesterification; ii) interesterification; iii) hydrolysis; iv) esterification; v) ammoniolysis; vi) aminolysis; and vii) perhydrolysis, 30 which method is characterized, in that a cross-linked lipase-aggregate is prepared according to the invention, and said cross-linked lipase-aggregate is contacted with a substrate.

The present invention will now be illustrated with reference to the following examples.

35

EXAMPLE 1.

Preparation of cross-linked lipases

- *R. miehei* lipase Lipozyme (Novo Nordisk)

- *Candida antarctica* lipase B SP525 (Novo Nordisk)
- *P. alcaligenes* lipase (Gist-brocades).

METHOD Ia) In the presence of (NH₄)₂SO₄ (Water)

0.5 ml stock enzyme solution (Lipozyme or SP525, Novo Nor-
5 disk, Copenhagen, Denmark) or 50 mg PaL enzyme powder (PaL is
P. alcaligenes lipase (Gist-Brocades, Delft, the Nether-
lands)), are dissolved in 1 ml potassium phosphate buffer
(100 mM, pH 7) in a 10 ml centrifuge tube. 550 mg (NH₄)₂SO₄
was added followed by 1 ml of a precipitating solution A con-
10 sisting of 550 mg (NH₄)₂SO₄ (the precipitant) in potassium
phosphate buffer (100 mM, pH 7), and 80 µl glutardialdehyde
(25% solution in water) are added. The mixture is left stir-
ring at 4°C (room temperature for SP525) for 17 hours.

3 ml H₂O is added to the mixture and then centrifuged. The su-
15 pernatant is decanted and the residue is washed, centrifuged
and decanted 3 more times with H₂O (5 ml each time). The final
enzyme preparation is kept in 5 ml H₂O. If necessary the solid
is dispersed by stirring with magnets.

METHOD Ib and Ic) In the presence of (NH₄)₂SO₄ and surfactant

20 The method Ia) was repeated, with the difference that the
precipitating solution A contained in addition 25 mg sodium
dodecylsulphate (SDS) (method Ib) or 25 mg Triton X-100 (TR)
(method Ic).

METHOD Id) In the presence of dimethoxyethane (DME)

25 0.5 ml stock enzyme solution (Lipozyme or SP525) or 50 mg PaL
enzyme powder, are introduced in a 10 ml centrifuge tube, af-
ter which 1 ml potassium phosphate buffer (100 mM, pH 7), 3
ml DME and 80 µl glutardialdehyde are added. The mixture is
left stirring at 4°C (room temperature for SP525) for 17
30 hours.

1 ml DME is added to the mixture and then centrifuged. The
supernatant is decanted and the residue is washed, centri-
fuged and decanted 3 more times with DME (5 ml each time).
The final enzyme preparation is kept in 5 ml DME. If neces-
35 sary the solid is dispersed by stirring with magnets.

METHOD Ie) In the presence of DME and a Crownether

Method Id) was repeated, except that an amount of 6.9 mg the
crownether dibenzo-18-crown-6 was added too together with the

3 ml DME.

Cross-linked lipase preparations were kept in suspension, either in water or DME, named preparations Ia-e.

5 EXAMPLE 2

Activity tests

The activity of CLEA's prepared according to the preparations Ia-e) were tested with three reactions. These reactions were performed either in an aqueous solution, or in an organic solution.
10

For tests IIb and IIc, the following general procedure was followed, before performing the assay:

- Of aqueous suspensions, 0.5 ml samples of CLEA suspension were lyophilized in 2 ml Eppendorf cup each.
- 15 - Of organic suspensions, 0.5 ml samples of CLEA suspension were centrifuged in a 2 ml Eppendorf cup each, and the supernatant was decanted.
- Native enzyme, used as a control, was used directly from the stock solution (50 µl), or as powder (5 mg), without further treatment.
20

For test IIb and IIc, analyses were performed using a GC Varian Star 3600 gas chromatograph (Varian). Column: CP WAX 52 CB 50 m x 0.53 mm. Carrier gas: nitrogen.

Test IIa) Hydrolysis of *p*-nitrophenyl propionate (performed
25 in water)

The enzyme preparation (50 µl) was suspended in 0.5 ml water (1.5 ml in case of SP525) and 200 µl (50 µl in case of SP525) of the suspension was added to 2.5 ml ester solution (0.4 mM *p*-nitrophenyl propionate in water at 25°C). Measurement of
30 the increase of absorbance at 348 nm (appearance of *p*-nitrophenol) in the hydrolysis of *p*-nitrophenyl propionate was monitored using a Cary 3-Bio UV spectrophotometer from Varian. Values are given as percentage of the activity of the native enzyme.

35 The result is summarized in table 1

Table 1

Hydrolysis of p-nitrophenyl propionate in water

Enz	Ia	Ib	Ic	Id	Ie	Native
Lipozyme	152	218	44	0	0	100
SP525	93	40	129	151	177	100

5 % of activity of native enzyme

Test IIb) Transesterification of ethyl octanoate with octanol (performed in diisopropylether)

- 10 0.5 ml diisopropyl ether is added to the Eppendorf cup containing the enzyme. 50 mg activated zeolite NaA (Aldrich) is added. 0.5 ml solution of 86 mM ethyl octanoate, 400 mM octanol, and 50 mM decane in diisopropyl ether is added and left stirring at room temperature. At the specified time, the reaction mixture is centrifuged, an aliquot (100 µl) withdrawn, dissolved in 0.5 ml DME, and analyzed by GC.
- 15 The result is summarized in table 2

Table 2

- 20 Transesterification of ethyl octanoate with octanol in i-Pr₂O

Enz	Ia	Id	Ic	Id	Ie	Native
Lipozyme	0.16	3.48	0.33	23.79	18.89	0.16
Pal	0.15	7.11	24.18	0.11	0.27	17.55

Activity expressed as µmol of octyl octanoate produced in 24 h.

- 25 *Test IIc) Hydrolysis of ethyl octanoate (performed in DME/water mixture)*
- 50 µl H₂O and 0.95 ml solution of 43 mM ethyl octanoate, and 25 mM decane is added to Eppendorf cup containing the enzyme
- 30 and left stirring at room temperature. At the specified time, anhydrous Na₂SO₄ is added to the reaction mixture, shaken, and centrifuged. An aliquot (100 µl) is then withdrawn, dissolved

in 0.5 ml DME, and analyzed by GC.
The result is summarized in table 3

Table 3

5

Hydrolysis of ethyl octanoate in DME/water

Enz	Ia	Ib	Ic	Id	Ie	Native
Lipo- zyme 24h	24.34	<u>40.48</u>	25.37	22.85	22.73	16.34
Pal. 3h	35.82	32.02	<u>40.26</u>	0.49	0.37	3.34

Activity expressed as μmol of octanoic acid produced in 24 or 3 h.

10

As shown by the above results, adding a surfactant or a crown ether in accordance with the present invention, may have a beneficial effect on the enzymatic activity.

15 EXAMPLE 3

Washing test

Methode Ia was repeated using SP525. The supernatant is decanted and the pellet resuspended in 5 ml water, centrifuged and the supernatant is decanted.

20 The pellet obtained is subjected to the same washing procedure with 3 * 5 ml water or DME. After the last washing step, the CLEA is kept in 5 ml of the same solvent.

The two CLEAs were tested using the transesterification of ethyl octanoate with octanol (as described in test IIc).

25 0.5 ml DME is added to 50 μl enzyme preparation in a 2 ml Eppendorf. The enzyme washed with water only was previously lyophilized in the Eppendorf cup. 50 mg activated zeolite NaA is added. 0.5 ml solution of 86 mM ethyl octanoate, 400 mM octanol, and 50 mM decane in DME is added and left stirring

30 at room temperature. After 1 hour, the reaction mixture is centrifuged, an aliquot (100 μl) withdrawn, dissolved in 0.5 ml DME, and analyzed by GC.

Table 4

Transesterification of ethyl octanoate with octanol (in DME)

5

Processing solvent	Activity
Water	5.41
DME	14.03

Activity expressed as μmol of octyl octanoate produced in 1 h.

10 EXAMPLE 4

Lipase CLEAs were prepared according the procedure described in example 1 and were tested in the hydrolysis of methyl mandelate. The enzyme was added to a 50 mM solution of methyl mandelate in dimethoxyethane-potassium phosphate buffer pH 7 (95:5, v/v). After 48 hours the conversion and enantiomeric excess of the substrate (ee_s) were analyzed using HPLC (see Table 5).

20

Table 5

Precipitant	Additive	Conv. (%)	ee_s	E
Dimethoxyethane	-	31.0	0.31 (R)	7
Dimethoxyethane	CR	33.5	0.33 (R)	7
Acetone	-	26.7	0.26 (R)	8
$(\text{NH}_4)_2\text{SO}_4$	SDS	27.9	0.27 (R)	8
$(\text{NH}_4)_2\text{SO}_4$	TR	23.2	0.24 (R)	10
Free enzyme	-	33.9	0.33 (R)	6

CR : Crownether dibenzo-18-crown-6

SDS: Sodium dodecylsulfate

TR : Triton X-100

ee_s : enantiomeric excess of the substrate

25 E : Enantiomeric ratio.

From table 5 it can be seen that the Enantiomeric ratio can be increased using the method according to the present invention.

CLAIMS

1. Method of preparing cross-linked enzyme aggregates comprising the steps of

a) precipitating a dissolved enzyme present in a liquid me-

5 dium;

b) cross-linking with a cross-linker, and

c) optionally, washing the cross-linked enzyme aggregate,

characterized, in that at least one treatment is performed chosen from the group consisting of I) performing step a) in

10 the presence of a compound selected from the group consisting

of i) a crown ether, and ii) a surfactant, and II) performing

step c) with a second liquid differing in organic composition from the liquid.

2. Method according to claim 1, **characterized**, in

15 that step a) and step b) are performed simultaneously.

3. Method according to claim 1 or 2, **characterized**,

in that the compound is removed after cross-linking.

4. Method according to any of the preceding claims, **characterized**, in that the enzyme is a lipase.

20 5. Method for performing a lipase-catalysed reac-

tion, said reaction being chosen from the group consisting of i) transesterification; ii) hydrolysis; and iii) ammoniolysis, **characterized**, in that a cross-linked lipase-aggregate

is prepared according to claim 4, and said cross-linked li-

25 pase-aggregate is contacted with a substrate.

INTERNATIONAL SEARCH REPORT

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A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 C12N9/96 C12N9/20

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 C12N C12P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, MEDLINE, PAJ, BIOSIS, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

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X	CAO L ET AL: "Cross-linked aggregates of penicillin acylase: Robust catalysts for the synthesis of beta-lactam antibiotics." JOURNAL OF MOLECULAR CATALYSIS B ENZYMATIC, vol. 11, no. 4-6, 2001, pages 665-670, XP001079507 ISSN: 1381-1177 the whole document ---	1
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Further documents are listed in the continuation of box C.

Patent family members are listed in annex.

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

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